AMENDMENTS TO THE CLAIMS

1. (Currently Amended) A compound represented by the formula (I):

$$\begin{array}{c} R^{7}OH \\ \downarrow \\ OH \end{array}$$

wherein R⁷ and R²¹ are the same or are different and represent

-O-benzoyl,

OH, or

RC(=Y)-O-, wherein Y represents an oxygen atom, and R represents

piperazinyl, alkyl, -O-phenyl, -N-alkyl or -NH-phenyl,

a C_6 to C_{14} aryl-group which may have a substituent, or

a C₆ to C₁₄ aryloxy group which may have a substituent, or

R^{N1}R^{N2}N R^M-, wherein R^M-represents

a) a single bond,

b) CO O,

c) -CS-O-or

d) CO NR^{N3}, wherein R^{N3} represents a hydrogen atom or a C₁ to C₆ alkyl group which may have a substituent, provided that, the leftmost bond in b) to e) is bonded to the nitrogen atom, and

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wherein R^{N1} and R^{N2} are the same or are different and represent

- a) a hydrogen atom,
- b) a C₁ to C₂₂ alkyl group which may have a substituent,
- e) an unsaturated C2 to C22 alkyl group which may have a substituent,
- d) an aliphatic C2 to C22 acyl group which may have a substituent,
- e) an aromatic C_7 to C_{15} acyl group which may have a substituent,
- f) a C₆ to C₁₄ aryl group which may have a substituent,
- g) a 5-membered to 14-membered heteroaryl group which may have a substituent,
- h) a C₇ to C₂₂ aralkyl group which may have a substituent,
- i) a C₁ to C₂₂ alkylsulfonyl group which may have a substituent,
- j) a C₆ to C₁₄ arylsulfonyl group which may have a substituent,
- k) a 3-membered to 14-membered non-aromatic heterocyclic group formed by R^{N1} and R^{N2} together in combination with the nitrogen atom to which R^{N1} and R^{N2} are bonded, wherein the 3-membered to 14-membered non-aromatic heterocyclic group may have a substituent,
- 1) a 5-membered to 14-membered heteroaralkyl group which may have a substituent,
- m) a C₃ to C₁₄ cycloalkyl group which may have a substituent or
- n) a 3-membered to 14-membered non-aromatic heterocyclic group which may have a substituent; or
 - a pharmacologically acceptable salt thereof, thereof.
- wherein said substituents are each independently selected from the group consisting of: C₁-C₆ alkyl group, phenyl group, halogen, hydroxyl group, C₁-C₆ alkoxy group, thiol group, C₁-

C₆-alkylthio group, nitro group, nitroso group, cyano group, C₁-C₆-alkoxycarbonyl group, amino group, mono (C₁-C₆-alkyl) amino group, di (C₁-C₆-alkyl) amino group, pyrrolidyl group, piperadyl group, piperidyl group and pyrridyl group.

2. (Currently Amended) The compound according to claim 1 represented by the formula (I-a):

wherein R^{7a} and R^{21a} are the same or are different and represent

 $R^aC(=Y^a)$ -O-, wherein Y^a represents an oxygen atom, and R^a represents

a C_6 to C_{14} aryl group which may have a substituent, or

a C₆ to C₁₄ aryloxy group which may have a substituent, or

 $R^{aN1}R^{aN2}N$ -CO-O-, wherein R^{aN1} and R^{aN2} , the same or different, represent

a) a hydrogen atom,

- [[b)]] a C₁ to C₂₂ alkyl group which may have a substituent,
- [[c)]] an unsaturated C₂ to C₂₂ alkyl group which may have a substituent,

d) a C₆ to C₁₄ aryl group which may have a substituent,

e) a 5-membered to 14-membered heteroaryl group which may have a substituent,

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f) a C7 to C22 aralkyl group which may have a substituent,

g) a 3-membered to 14-membered non-aromatic heterocyclic group formed by R^{aN1} and R^{aN2} together in combination with the nitrogen atom to which R^{aN1} and R^{aN2} are bonded, wherein the 3-membered to 14-membered non-aromatic heterocyclic group may have a substituent,

h) a 5-membered to 14-membered heteroaralkyl group which may have a substituent,

[[i)]] a C₃ to C₁₄ cycloalkyl group which may have a substituent or

j) a 3-membered to 14-membered non-aromatic heterocyclic group which may
have a substituent, or

R^{aN1}R^{aN2}N-CS-O-, wherein R^{aN1} and R^{aN2} are the same as defined above,

[[;]] or a pharmacologically acceptable salt thereof, thereof.

wherein said substituents are each independently selected from the group consisting of: C₁-C₆ alkyl group, phenyl group, halogen, hydroxyl group, C₁-C₆ alkoxy group, thiol group, C₁-C₆ alkylthio group, nitro group, nitroso group, cyano group, C₁-C₆ alkoxycarbonyl group, amino group, mono (C₁-C₆ alkyl) amino group, di (C₁-C₆ alkyl) amino group, pyrrolidyl group, piperadyl group, piperidyl group and pyrridyl group.

- 3. (Canceled).
- 4. (Currently Amended) The compound according to claim 1, wherein R^7 and R^{21} R^{N+1} and R^{N+2} are the same or are different and represent a C_1 to C_6 alkyl group or C_6 to C_{14} aryl group,

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or form, together in combination with the nitrogen atom to which R^{NI} and R^{N2} are bonded, a non-aromatic heterocyclic group selected from the group consisting of:

or a pharmacologically acceptable salt thereof.

5-18. (Canceled).

19. (Currently Amended) The compound according to claim 1, which is (8E,12E,14E)-21-benzoyloxy-3,6-dihydroxy-6,10,12,16,20-pentamethyl-7-((4-methylpiperazin-1-yl)carbonyl)oxy-18,19-epoxytricosa-8,12,14-trien-11-olide,

(8E,12E,14E)-21-(N,N-dimethylcarbamoyloxy)-3,6-dihydroxy-6,10,12,16,20-pentamethyl-7((4-methylpiperazin-1-yl)carbonyl)oxy-18,19-epoxytricosa-8,12,14-trien-11-olide, (8E,12E,14E)-3,6-dihydroxy-6,10,12,16,20-pentamethyl-21-N,N-dimethylcarbamoyloxy-7 ((4-methylpiperazin-1-yl)carbonyl)oxy-18,19-epoxytricosa-8,12,14-trien-11-olide and (8E,12E,14E)-3,6-dihydroxy-6,10,12,16,20-pentamethyl-7-((4-methylpiperazin-1-yl)carbonyl)oxy-21-phenylcarbamoyloxy-18,19-epoxytricosa-8,12,14-trien-11-olide; or a

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pharmacologically acceptable salt thereof.

20. (Canceled).

21. (Previously Presented) A pharmaceutical composition comprising the compound

according to claim 1, or a pharmacologically acceptable salt thereof as an active ingredient and a

pharmaceutically acceptable carrier.

22-45. (Canceled).

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